



Post Authorisation Assessments

Leventia 1 mg/ml Oral Solution for Dogs Vm 06376/3058

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| 02 September 2025 | Change of legal entity of the Marketing Authorisation Holder from MSD Animal Health UK Limited, Walton Manor, Walton, Milton Keynes, MK7 7AJ, United Kingdom to Intervet International B.V., Wim de Körverstraat 35, 5831 AN Boxmeer, Netherlands. |
| 24 July 2025 | One-off alignment of the product information with version 9.0* of the QRD templates. |
| 14 December 2024 | Update to a Ph. Eur. CEP for an already authorised manufacturer of the active substance. (NI) |
| 14 December 2024 | Update to a Ph. Eur. CEP for an already authorised manufacturer of the active substance. (GB) |
| 26 April 2024 | Change in the dimensions of the immediate packaging for the finished product. (NI) |
| 25 January 2024 | Alternative primary packaging container registered for the finished product. (GB) |
| 25 August 2021 | Addition of a site where batch control/testing takes place. Addition of a site where batch control/testing takes place. |
| 27 April 2021 | Submission of an updated Ph. Eur. certificate of suitability for an active substance from an already approved manufacturer. |
| 14 August 2020 | Change in the name of the marketing authorisation holder from Intervet UK Limited to MSD Animal Health UK Limited. |
| 13 November 2018 | Change in the safety database of an existing pharmacovigilance system as described in the DDPS |
| 16 January 2018 | Change in the specification limits of the finished product. |
| 01 December 2016 | Change in the safety database of an existing pharmacovigilance system as described in the DDPS. |
| 27 April 2016 | Submission of an updated Ph. Eur. certificate of suitability. |
| 02 April 2014 | Reduction in the shelf-life of the finished product, from 3 years to 2 years. |
| 27 November 2014 | Update of the pharmacovigilance system as described in the DDPS. |
| 06 October 2014 | Change in specification limit of the finished product. |
| 22 October 2012 | Changes to the DDPS following assessment of the same DDPS in relation to another medicinal product of the same MAH. |
| 13 June 2012 | Renewal procedure – Ireland as RMS. |
| 18 March 2009 | Change shelf life of finished product (as packaged for sale) |
| 19 November 2008 | Simple dosage instruction changes |
| 22 May 2008 | Change shelf life of finished product (after first opening) |

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| 22 May 2008 | Formulation |
| 04 July 2007 | Deletion of any manufacturing site |